LISTING OF CLAIMS

1-4. (Canceled)

5. (Currently Amended) The method according to claim 1 wherein the nucleic acid molecule and the prokaryotic DNA ligase polypeptide are contacted A method of modifying a nucleic acid molecule comprising:

contacting the nucleic acid molecule with an isolated prokaryotic DNA ligase polypeptide in the presence of a prokaryotic Ku polypeptide, wherein the prokaryotic DNA ligase polypeptide has at least 95% amino acid sequence identity with the amino acid sequence of SEQ ID NO: 91 and the prokaryotic Ku polypeptide shares has at least 95% amino acid sequence identity with the amino acid sequence of SEQ ID NO: 92 accession number CAB08491 (SEQ ID NO: 92).

6-7. (Canceled)

8. (Currently Amended) A method of ligating nucleic acid molecule ends comprising:

contacting a first nucleic acid end and a second nucleic acid end with an isolated prokaryotic DNA ligase polypeptide,

wherein said first and said second nucleic acid ends comprise non-complementary overhang regions, and

wherein the prokaryotic DNA ligase polypeptide shares has at least 95% amino acid sequence identity with the amino acid sequence of <u>SEQ ID NO: 91</u>-accession-number CAB08492 (SEQ ID NO: 91).

9. (Canceled)

10. (Previously Presented) The method according to claim 8 wherein the first end is on a first nucleic acid molecule and the second end is on a second nucleic acid molecule.

- 11. (Previously Presented) The method according to claim 10 wherein the first and second nucleic acid molecules are DNA.
- 12. (Previously Presented) The method according to claim 10 wherein the first nucleic acid molecule is DNA and the second nucleic acid molecule is RNA.
- 13. (Previously Presented) The method according to claim 8 wherein the first and second ends are on the same nucleic acid molecule.
- 14. (Previously Presented) The method according to claim 8 further comprising isolating the ligated nucleic acid molecule, purifying the ligated nucleic acid molecule, or both isolating and purifying the ligated nucleic acid molecule.
- 15. (Withdrawn and Currently Amended) A method of labeling a nucleic acid molecule comprising:

contacting a nucleic acid molecule having a first terminus with an isolated prokaryotic DNA ligase polypeptide in the presence of labelled nucleotides, wherein the prokaryotic DNA ligase polypeptide shares has at least 95% amino acid sequence identity with the amino acid sequence of SEQ ID NO: 91-accession number CAB08492 (SEQ ID NO: 91).

- 16. (Withdrawn) The method according to claim 15 wherein the nucleotides are NTPs.
- 17. (Withdrawn) The method according to claim 15 wherein the nucleotides are dNTPs.
- 18. (Currently Amended) A method of filling in a single stranded gap in a double stranded nucleic acid molecule comprising:

contacting a double stranded nucleic acid molecule having a single stranded region with an isolated prokaryotic DNA ligase polypeptide, wherein the prokaryotic DNA ligase

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polypeptide shares has at least 95% amino acid sequence identity with the amino acid sequence of <u>SEQ ID NO: 91</u> accession number CAB08492 (SEQ ID NO: 91).

- 19. (Previously Presented) The method according to claim 18 wherein said nucleic acid molecule and said prokaryotic DNA repair ligase polypeptide are contacted in the presence of NTPs.
- 20. (Previously Presented) The method according to claim 18 wherein said nucleic acid molecule and said prokaryotic DNA repair ligase polypeptide are contacted in the presence of dNTPs.
- 21. (Currently Amended) A method of removing a single stranded overhang from the end of a nucleic acid molecule comprising:

contacting said nucleic acid molecule with an isolated prokaryotic DNA ligase polypeptide, wherein the prokaryotic DNA ligase polypeptide shares has at least 95% sequence identity with the amino acid sequence of <u>SEQ ID NO: 91</u> accession number CAB08492 (SEQ ID NO: 91).

22. (Canceled)

- 23. (Previously Presented) The method according to claim 21 wherein said nucleic acid molecule is contacted in the presence of Mg²⁺ or Mn²⁺.
- 24. (Withdrawn and Currently Amended) A method of producing an RNA molecule comprising:

contacting an isolated prokaryotic DNA ligase polypeptide and a template DNA strand in the presence of NTPs, wherein the prokaryotic DNA ligase polypeptide shares-has at least 95% amino acid sequence identity with the amino acid sequence of SEQ ID NO: 91-accession number CAB08492 (SEQ ID NO: 91).

- 25. (Withdrawn) The method according to claim 24 wherein the isolated prokaryotic DNA ligase and the template DNA are contacted in the presence of a primer oligonucleotide.
- 26. (Withdrawn and Currently Amended) A method of producing an DNA molecule comprising:

contacting an isolated prokaryotic DNA ligase polypeptide and a nucleic acid template in the presence of dNTPs and a primer oligonucleotide, wherein the prokaryotic DNA ligase polypeptide shares has at least 95% amino acid sequence identity with the amino acid sequence of SEQ ID NO: 91 accession number CAB08492 (SEQ ID NO: 91).

27. (Withdrawn) The method according to claim 26 wherein the nucleic acid template is an RNA template.

28-30. (Canceled)

31. (Currently Amended) The method according to claim 8 wherein the nucleic acid molecule and the prokaryotic DNA ligase polypeptide are contacted in the presence of a prokaryotic Ku polypeptide, wherein the prokaryotic Ku polypeptide shares has at least 95% amino acid sequence identity with the amino acid sequence of SEQ ID NO: 92-accession number CAB08491 (SEQ ID NO: 92).

32-33. (Canceled)

- 34. (Withdrawn and Currently Amended) A kit comprising an isolated prokaryotic DNA ligase polypeptide for use in a method according to claim 1, wherein the isolated prokaryotic DNA ligase polypeptide shares has at least 95% amino acid sequence identity with the amino acid sequence of SEQ ID NO: 91 accession number CAB08492 (SEQ ID NO: 91).
- 35. (Withdrawn and Currently Amended) The kit according to claim 34 comprising an isolated Mt-Ku polypeptide, wherein the prokaryotic Ku polypeptide shares-has at least 95%

amino acid sequence identity with the amino acid sequence of <u>SEQ ID NO: 92-accession number</u> CAB08491 (SEQ ID NO: 92).

- 36. (Withdrawn) The kit according to claim 34 comprising dNTPs.
- 37. (Withdrawn) The kit according to claim 34 comprising NTPs.
- 38. (Withdrawn) The kit according to claim 34 comprising one or more of buffers, stabilisers and excipients.
- 39. (Withdrawn) A method of producing a prokaryotic DNA repair polypeptide comprising:
- (a) causing expression from a nucleic acid which encodes a prokaryotic DNA repair polypeptide in a suitable expression system to produce the polypeptide recombinantly; and,
- (b) testing the recombinantly produced polypeptide for prokaryotic DNA repair activity.
- 40. (Withdrawn) The method according to claim 39 wherein the recombinantly produced polypeptide is tested for one or more of: non-complementary end ligation activity, DNA dependent RNA primase activity, 3'-5' exonuclease activity, DNA and RNA dependent DNA polymerase activity, DNA dependent RNA polymerase activity, ATP dependent DNA and RNA ligase activity and DNA terminal transferase activity.
- 41. (Withdrawn) The method according to claim 39 wherein the prokaryotic DNA repair polypeptide is an Mt-Lig polypeptide or an allele or variant thereof.
- 42. (Withdrawn) The method according to claim 39 comprising purifying said recombinantly produced polypeptide.
- 43. (Withdrawn) The method according to claim 26 wherein the nucleic acid template is a DNA template.

- 44. (New) The method according to claim 18 wherein the double stranded nucleic acid molecule and the prokaryotic DNA ligase polypeptide are contacted in the presence of a prokaryotic Ku polypeptide, wherein the prokaryotic Ku polypeptide has at least 95% amino acid sequence identity with the amino acid sequence of SEQ ID NO: 92.
- 45. (New) The method according to claim 21 wherein the nucleic acid molecule and the prokaryotic DNA ligase polypeptide are contacted in the presence of a prokaryotic Ku polypeptide, wherein the prokaryotic Ku polypeptide has at least 95% amino acid sequence identity with the amino acid sequence of SEQ ID NO: 92.